		1623	RECEIVED VTRAL FAX CENTER	
Response To OFFICIAL ACTION				PAGE NUMBER
Respo	Krishnan, Ganapathy			
10/736.084 12/15/2003 Joseph C. Walsh			2003P88074US EXAMINER	
APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNE	Y DOCKET NO.

AMENDMENTS TO THE CLAIMS

In the Claims, please make the following amendments:

1. (Previously presented) A method for preparing a compound having the following formula:

wherein R is an alkoxy blocking group; P is a hydroxyl protecting group; and L is a leaving group, the method comprising the steps of:

reacting a compound of the formula:

with a hydroxyl protecting group to produce a compound having the following formula:

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNE	Y DOCKET NO.
10/736,084 12/15/2003 Joseph C. Walsh			2003P88074US EXAMINER Krishnan, Ganapathy	
Respo		,		
·		ART UNIT	PAGE NUMBER	
			1623	9

wherein P is the same as defined above;

b. enolating the reaction product of step (a) produce a compound having the following formula:

wherein P and R are the same as defined above; and

c. incorporating a leaving group to produce a compound having the following formula:

2. (Original) The method according to Claim 1, wherein P is selected from the group consisting of methoxymethyl ether, methylthiomethyl ether, 2-methoxyethoxymethyl ether, 1-ethoxyethyl ether, 1-methyl-1-methoxyethyl ether, t-butyl ether, allyl ether, benzyl ether, 4-nitrobenzyl ether, o-nitrobenzyl ether, trityl ether, monomethoxytrityl ether, dimethoxytrityl ether, tritylone ether, tetrahydropyran ether, tetrahydropyranyl ether, 4-methoxy tetrahydropyran ether, 4-methoxytetrahydrothiopyranyl ether, tetrahydrofuran ether, tetrahydrotriofuranyl ether, isobutyrate ester, pivaloate ester, adamantoate ester, benzoate ester, 2,4,6,-trimethylbenzoate ester, methyl carbonate,

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNE	EY DOCKET NO.
10/736,084	12/15/2003	Joseph C. Walsh	2003P88074US	
	EXAMINER Krishnan, Ganapathy			
Respo				
1100			ART UNIT	PAGE NUMBER
-			1623	

allyl carbonate, benzyl carbonate, p-nitrobenzyl carbonate, t-Bu carbonate, S-benzylthio carbonate, N-phenyl carbamate, and nitrate ester.

- 3. (Original) The method according to Claim 1, wherein P is selected from the group consisting of dimethoxytrityl, monomethoxytrityl, trityl, t-butyloxycarbonyl, t-butyldimethylsilyl, t-butyldiphenylsilyl, tetrahydropyranyl ether, tetrahydrofuranyl ether, ethoxyethyl ether, and 1-methyl-1-methoxyethyl ether.
- 4. (Original) The method according to Claim 1, wherein R is alkyl C_1 - C_4 , *i*-propyl, benzyl, cycloalkane C_3 - C_6 , phenyl, tosyl, acetate, or benzoate.
- 5. (Original) The method according to Claim 1, wherein R is methyl, ethyl, *i*-propyl, benzyl, or cycloalkane C₃-C ₆.
- 6. (Original) The method according to Claim 1, wherein step (b) includes treating the reaction product of step (a) with an alkoxide having 1 to 4 carbons, cycloalkoxide C₃-C₆, phenoxide, tosyate, acetate, or benzoate.
- 7. (Original) The method according to Claim 6, wherein the alkoxide is sodium methoxide.
- 8. (Original) The method according to Claim 1, wherein L is a sulfonate ester.
- 9. (Original) The method according to Claim 1, wherein L is selected from the group consisting of mesylate, nosylate, tosylate, and triflate.
- 10. (Previously presented) A method for preparing a precursor for the preparation of a radiolabeled nucleoside comprising:
 - a. converting a 2-deoxy nucleoside into a 2,3'-anhydronucleoside;
- b. reacting the 2,3'-anhydronucleoside with a hydroxyl protecting group to produce a 2,3'-anhydronucleoside derivative wherein the 5'-O group is protected;

APPLICATION NO. FILING DATE FIRST NAMED INVE			ATTORNEY DOCKET NO.		
10/736,084 12/15/2003 Joseph C. Walsh		2003P88074US			
	•.	EXAMINER			
Respo	Krishnan, Ganapathy				
		ART UNIT	PAGE NUMBER		
			1623		

- c. reacting the protected 2,3'-anhydronucleoside derivative with a reagent that opens the 2,3'-anhydro-ring and enolates the 2-position on the pyrimidine ring; and
- d. incorporating a leaving group to produce the radiolabeled nucleoside precursor;

where the nucleoside base is thymidine or uridine.

- 11. (Previously presented) The method according to Claim 10, wherein the nucleoside is thymidine or uridine.
- 12. (Original) A method for preparing a precursor for the preparation of 3'-Deoxy-3'-[¹⁸F]-fluoro-thymidine (¹⁸F-FLT) comprising:
 - a. converting thymidine into 2,3'-anhydrothymidine;
- b. reacting the 2,3'-anhydro thymidine with a hydroxyl protecting group to produce a 2,3'-anhydrothymidine derivative wherein the 5'-O group is protected;
- c. reacting the protected 2,3'-anhydrothymidine derivative with a reagent that opens the 2,3'-anhydro-ring and enolates the 2-position on the pyrimidine ring; and
 - d. incorporating a leaving group to produce the ¹⁸F-FLT precursor.
- 13. (Original) The method according to Claim 12, wherein step (c) produces an enol having an -O-R group attached to the 2-carbon.
- 14. (Previously presented) A method according to Claim 13, wherein R is alkyl C₁-C₄, benzyl, cycloalkane C₃-C₆, phenyl, tosyl, acetate, or benzoate.
- 15. (Original) A method according to Claim 12, wherein step (c) includes treating the reaction product of step (b) with an alkoxide.
- 16. (Previously presented) A method according to Claim 15, wherein the alkoxide is selected from the group consisting of sodium methoxide, and sodium ethoxide.

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNE	EY DOCKET NO.	
10/736,084	10/736,084 12/15/2003 Joseph C. Walsh			2003P88074US	
	EXAMINER Krishnan, Ganapathy				
Respo					
		ART UNIT	PAGE NUMBER		
			1623	P	

- 17. (Original) A method according to Claim 12, wherein the hydroxyl protecting group is dimethoxytrityl, monomethoxytrityl, trityl, t-butyloxycarbonyl, t-butyldimethylsilyl, t-butyldiphenylsilyl, tetrahydropyranyl ether, tetrahydrofuranyl ether, ethoxyethyl ether, or 1-methyl-1-methoxyethyl ether.
- 18. (Original) A method according to Claim 12, wherein the hydroxyl protecting group is dimethoxytrityl, monomethoxytrityl, or trityl.
- 19. (Original) A method according to Claim 12 wherein the leaving group is a sulfonate ester.
- 20. (Original) A method according to Claim 19, wherein the leaving group is mesylate, tosylate, nosylate, or triflate.
- 21. (Previously presented) A compound having the following formula:

wherein R is alkyl C_1 - C_4 , benzyl, cycloalkane C_3 - C_6 , phenyl, tosyl, acetate, or benzoate; P is a hydroxyl protecting group; and L is a leaving group.

- 22. (Original) A compound according to Claim 21, wherein R is methyl or ethyl.
- 23. (Original) A compound according to Claim 21, wherein P is methoxymethyl ether, methylthiomethyl ether, 2-methoxyethoxymethyl ether, 1-ethoxyethyl ether, 1-methyl-1-methoxyethyl ether, t-butyl ether, allyl ether, benzyl ether, 4-nitrobenzyl ether, o-nitrobenzyl ether, trityl ether, monomethoxytrityl ether, dimethoxytrityl ether, tritylone ether; tetrahydropyran ether, tetrahydrothiopyranyl ether, 4-methoxy tetrahydropyran ether, 4-methoxytetrahydrothiopyranyl ether, tetrahydrofuran ether, tetrahydrotriofuranyl

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNE	Y DOCKET NO.
10/735,084	12/15/2003	Joseph C. Walsh	2003P88074US	
	EXAMINER			
Resno	Krishna	n, Ganapathy		
Response To OFFICIAL ACTION			ART UNIT	PAGE NUMBER
			1623	_

ether, isobutyrate ester, pivaloate ester, adamantoate ester, benzoate ester, 2,4,6,-trimethylbenzoate ester; methyl carbonate, allyl carbonate, benzyl carbonate, p-nitrobenzyl carbonate, t-Bu carbonate, S-benzylthio carbonate, N-phenyl carbamate, or nitrate ester.

24. (Original) A compound according to Claim 21, wherein P is dimethoxytrityl, monomethoxytrityl, trityl, t-butyloxycarbonyl, t-butyldimethylsilyl, t-butyldiphenylsilyl, tetrahydropyranyl ether, tetrahydrofuranyl ether, ethoxyethyl ether, or 1-methyl-1-methoxyethyl ether.

25. (Original) A compound according to Claim 21, wherein P is dimethoxytrityl.

26. (Original) A compound according to Claim 21, wherein L is a sulfonate ester.

27. (Original) A compound according to Claim 21, wherein L is selected from the group consisting of p-(2,4-dinitroanilino)benzenesulfonyl, benzenesulfonyl, methylsulfonyl (mesylate), p-methylbenzenesulfonyl (tosylate), 4-nitrobenzene sulfonyl (nosylate), p-bromobenzenesulfonyl, trifluoromethylsulfonyl (triflate), trichloroacetimidate, acyloxy, 2,2,2-trifluoroethanesulfonyl, imidazolesulfonyl, and 2,4,6-trichlorophenyl.

28. (Original) A compound according to Claim 21, wherein R is methyl, P is dimethoxy trityl, and L is mesylate, tosylate, or nosylate.

29. (Original) A compound having the following formula:

wherein Ms is methylsulfonyl.

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNE	Y DOCKET NO.
10/736,084 12/15/2003 Joseph C. Walsh			2003P88074US	
		EXAMINER		
Respo	onse To OFFICIA I	Krishne	n, Genepathy	
			ART UNIT	PAGE NUMBER
			1623	

30. (Previously presented)

A compound having the following formula:

wherein R is alkyl C₁-C₄, benzyl, cycloalkane C₃-C₆, phenyl, tosyl, acetate, or benzoate; P is a hydroxyl protecting group; X is oxygen, sulfur, or nitrogen, and L is a leaving group.

31. (Original) A compound according to Claim 30, wherein L is halogen, p-(2,4-dinitroanilino)benzenesulfonyl, benzenesulfonyl, methylsulfonyl (mesylate), p-methylbenzenesulfonyl (tosylate), 4-nitrobenzene sulfonyl (nosylate), p-bromobenzenesulfonyl, trifluoromethylsulfonyl (triflate), trichloroacetimidate, acyloxy, 2,2,2-trifluoroethanesulfonyl, imidazolesulfonyl, or 2,4,6-trichlorophenyl.

32. (Original) A compound according to Claim 30, wherein P is dimethoxytrityl, monomethoxytrityl, trityl, t-butyloxycarbonyl, t-butyldimethylsilyl, t-butyldiphenylsilyl, tetrahydropyranyl ether, tetrahydrofuranyl ether, ethoxyethyl ether, or 1-methyl-1-methoxyethyl ether.

33-34. (Canceled)